Remove

Under the Paperwork Reduction Act of 1995, no persons are required to re

INFORMATION DISCLOSURE
STATEMENT BY APPLICANT
( Not for submission under 37 CFR 1.99)
(·,

Application Number		10562875		
Filing Date		2005-12-29		
First Named Inventor Mark		s Neumann, et al		
Art Unit		2681		
Examiner Name				
Attorney Docket Number		DE030230		

				0.3.			memore			
Examiner   Cite   Patent Number   Kind   Code		Issue Date  Name of Patentee or Applicant of cited Document		Pages,Columns,Lines where Relevant Passages or Relevan Figures Appear						
	1	6148187	A	2000-11-14	CHIBA					
	2	6278867	B1	2001-08-21	NORTHOUT	T, ET AL				
If you wisi	h to a	dd additional U.S. Pate	nt citatio	n information p	lease click the	Add button.	_	Add		
			U.S.P	ATENT APPLI	CATION PUB	LICATIONS		Remove		
Examiner Initial*	Cite Number Kind Code Date Name of Patentee or Applican of cited Document		Releva	Columns,Lines where int Passages or Relev s Appear						
	1	dd additional U.S. Pub						Add		
ir you wisi	n to a	dd additional U.S. Pub		FORFIGN PA			a button	Remove		
						MENTS		rtemove		
Examiner Initial*	Cite No	Foreign Document Number <sup>3</sup>	Country Code <sup>2</sup>	Kind	Publication	Name of Patente Applicant of cited Document	e or	Pages,Columns,Lines where Relevant Passages or Relevant Figures Appear		
Examiner Initial*				Kind	Publication	Name of Patente Applicant of cited	e or	Pages,Columns,Lines where Relevant Passages or Relevant		

U.S. PATENTS

## INFORMATION DISCLOSURE STATEMENT BY APPLICANT A ( Not for submission under 37 CFR 1.99) F

Application Number		10562875		
Filing Date		2005-12-29		
First Named Inventor	Mark.	us Neumann, et al		
Art Unit		2681		
Examiner Name				
Attorney Docket Number		DE030230		

	3	WO03032516		A1	2003-04-17	MOTOROLA INC		
	4	GB2354403		A	2001-03-21	ERICSSON		
	5	GB2317281		A	1998-03-18	MOTOROLA LTD		
If you wish to add additional Foreign Patent Document citation information please click the Add button Add								
NON-PATENT LITERATURE DOCUMENTS Remove								
Examiner	Cite	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item						

(book, magazine, journal, serial, symposium, catalog, etc), date, pages(s), volume-issue number(s),

If you wish to add additional non-patent literature document citation information please click the Add button Add

## EXAMINER SIGNATURE

publisher, city and/or country where published.

Examiner Signature Date Considered

\*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through a citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

1 See Kind Codes of USPTO Patent Documents at www.USPTO.GOV or MPEP 901.04. 2 Enter office that issued the document, by the two-letter code (WIPO Standard ST.3), 3 For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST.16 if possible. 5 Applicant is to place a check mark here if English language translation is attached.

Initials\* Nο

# INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

Application Number		10562875
Filing Date		2005-12-29
First Named Inventor Marks		us Neumann, et al
Art Unit		2681
Examiner Name		
Attorney Docket Numb	ec	DE030230

#### CERTIFICATION STATEMENT

Diagra can	37	CFR .	1 97	and	1 02 1	n make	the	appropriate	coloction/	e١٠

That each item of information contained in the information disclosure statement was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of the information disclosure statement. Sea 97 CFF 1.97(e)(1).

### OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquity, no item of information contained in the information disclosure statement was known to any individual designated in 37 CFR 1.56(c) more than three months prior to the filing of the information disclosure statement. See 37 CFR 1.97(c)(c)

- See attached certification statement.
- Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.

#### SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

Signature	/DAVID L. BARNES/	Date (YYYY-MM-DD)	2006-10-16
Name/Print	David Rames	Registration Number	47407

This collection of information is required by 3T CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is for lie fand by the USPTO to process) an application. Confidentiality is governed by \$5.1.S.C. 12.04 and 3T CFR 1.14. This collection is estimated to take 1 hour to complete, including gathering, preparing and submitting the completed application from the USPTO. Time will vary depending upon the individual case: Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. operatment of Commence; P.O. Box 1450, Alexandria, V.S. 231-1450, D.NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, V.S. 2211-1450.

### Privacy Act Statement

The Privacy Act of 1974 (P. L. 93-579) requires that you be given certain information in connection with your submission of the stackhold from related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, places be advised that (1) the general authority for the collection of this information is 35 U.S.C. 2(b)(2); (2) familishing of the information solicided is couldrain; and (3) the primoral pursuance for which the information is used by the U.S. Patient and Trademan Coffice is to process and/or examine your submission related to a patient agricultant or patient. If you do not furnish the requested process and/or examine your submission related to a patient agricultant or patient. If you do not furnish the requested results of the patient of the patient and the patient of the patient

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
  - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement negotiation.
  - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record pertains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
  - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974, as amended, pursuant to 5 U.S.C. 552(m).
  - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
    may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
    to the Patent Cooperation Treaty.
  - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
  - 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designed, uturing an insection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 4d U.S.C. 2904 and 2905. Such disclosure shall be made in accordance with the GSA requisions governing inseption of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make determinations about individuals.
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the
  application pursuant to 35 U.S.C. 12(2) to rissuance of a patent pursuant to 35 U.S.C. 151. Further, a record may be
  disclosed, subject to the limitations of 37 CFR 1.14, as a routine use, to the public if the record was filled in application
  which became abandoned or in which the proceedings were terminated and which application is referenced by either a
  published application, an application open to public inspections or as issued patent.
  - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.